I	H.28
2	Introduced by Representatives Till of Jericho, Brumsted of Shelburne,
3	Colburn of Burlington, Conlon of Cornwall, Conquest of
4	Newbury, Donovan of Burlington, Houghton of Essex, Lanphe
5	of Vergennes, Macaig of Williston, McCullough of Williston,
6	Mrowicki of Putney, Ode of Burlington, Patt of Worcester,
7	Rachelson of Burlington, Squirrell of Underhill, Sullivan of
8	Burlington, Troiano of Stannard, Yacovone of Morristown, and
9	Yantachka of Charlotte
10	Referred to Committee on
11	Date:
12	Subject: Health; regulated drugs; pharmacist; proper disposal
13	Statement of purpose of bill as introduced: This bill proposes to require
14	pharmacists dispensing regulated drugs to provide patients with information
15	regarding the importance of and processes for proper disposal of unused
16	medication.
17	An act relating to pharmacists providing information on the proper disposal
18	of unused regulated drugs

1	It is hereby enacted by the General Assembly of the State of Vermont:
2	Sec. 1. 18 V.S.A. § 4215 is amended to read:
3	§ 4215. AUTHORIZED SALES BY PHARMACISTS
4	* * *
5	(d) When dispensing a regulated drug pursuant to this section, a pharmacist
6	shall provide to the patient written information regarding the importance of
7	safely disposing of unused regulated drugs and how to do so properly.
8	Sec. 2. BOARD OF PHARMACY; PROPER DISPOSAL; RULEMAKING
9	The Board of Pharmacy, in consultation with the Department of Health,
10	shall adopt rules pursuant to 3 V.S.A. chapter 25 to establish the format and
11	content of the written information that pharmacists shall provide to patients
12	pursuant to 18 V.S.A. § 4215 on the importance of and processes for proper
13	disposal of unused regulated drugs.
14	Sec. 3. EFFECTIVE DATES
15	(a) Sec. 1 (18 V.S.A. § 4215) shall take effect on July 1, 2019.
16	(b) Sec. 2 (Board of Pharmacy rulemaking) and this section shall take
17	effect on passage, provided that the Board of Pharmacy shall provide
18	pharmacists with a reasonable period of time following adoption of the Board's
19	rules to make any necessary changes to their patient information materials to

ensure compliance with the rules' format and content specifications.

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